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Equip F vet. Injektionsvätska, suspension

Authorised

- Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated

Product identification

Medicine name:
Equip F vet. Injektionsvätska, suspension

Active substance:
Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
1.20 log10 haemagglutination inhibiting unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
2.10 log10 haemagglutination inhibiting unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated
2.40 log10 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis Animal Health ApS

Marketing authorisation date:
22/08/2003

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
18954

Date of authorisation status change:
22/08/2003

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.